Amendments to the Claims

This listing of claims replaces all other listings of claims:

1 - 9. (CANCELED)

10. (CURRENTLY AMENDED) A therapeutic method comprising providing to an eye of a patient an ecular a physiologic ophthalmic imigating or volume replacement solution containing at least one of a macrolide antibiotic or mycophenolic acid at a concentration in the range between about 1 ng/ml to about 200 µg/ml to provide irrigation, wash, or volume replacement further providing an anti-inflammatory effect without increased intraocular pressure a therapeutic effect.

11. (ORIGINAL) The method of claim 10 wherein the macrolide antibiotic or mycophenolic acid further provides at least one of an anti-inflammatory effect, an anti-cell proliferation effect, an anti-cell migration effect, an anti-angiogenesis effect, an antimicrobial effect, and an antifungal effect.

12, (CANCELED)

13. (CURRENTLY AMENDED) The method of claim 10 wherein the macrolide antibiotic or mycophenolic acid <u>further</u> provides an anti-angiogenic effect in a patient with an ocular tumor, a patient with diabetes, or a patient with sickle cell anemia.

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- 14. (ORIGINAL) The method of claim 10 wherein the macrolide antibiotic or mycophenolic acid is at a concentration of about 1 μg/ml.
- 15. (ORIGINAL) The method of claim 10 wherein the macrolide antibiotic or mycophenolic acid is at a concentration ranging from about 1 ng/ml to about 20 μg/ml.
- 16. (ORIGINAL) The method of claim 10 wherein the macrolide antibiotic or mycophenolic acid is at a concentration ranging from about 20 μg/ml to about 200 μg/ml.
- 17. (ORIGINAL) A therapeutic method comprising intraocularly administering to a patient undergoing cataract surgery an ocular solution containing at least one of a macrolide antibiotic or mycophenolic acid at a concentration in the range from about 20 μg/ml to about 200 μg/ml within a lens capsule prior to insertion of a replacement intraocular lens.
- 18. (ORIGINAL) The method of claim 17 wherein the solution reduces opacification of the posterior capsule.
- 19. (ORIGINAL) The method of claim 17 wherein the macrolide antibiotic is formulated as at least one of a liposome, a macrosphere, a microsphere, a macrocapsule, a microcapsule, a macrovesicle, and a microvesicle.

- 20. (ORIGINAL) The method of claim 17 wherein the macrolide antibiotic or mycophenolic acid is at a concentration in the range of about 20 μg/ml to about 200 μg/ml.
- 21. (ORIGINAL) The method of claim 19 wherein the macrolide antiblotic or mycophenolic acid is implanted within the capsule.
- 22. (CURRENTLY AMENDED) An article comprising an implantable ocular replacement lens in a solution containing a concentration of a macrolide antibiotic or mycophenolic acid ranging from about 20 µg/ml to about 2000 µg/ml sufficient to provide the lens with at least one effect selected from anti-cell proliferation, anti-cell migration, anti-inflammatory, anti-angiogenesis, antimicrobial, and antifungal.
- 23. (CANCELED)
- 24. (ORIGINAL) The article of claim 22 wherein the concentration is the range between about 20 μg/ml to about 200 μg/ml.
- 25. (ORIGINAL) The article of claim 22 wherein the macrolide antibiotic is at least one of tacrolimus, cyclosporine, sirolimus, everolimus, ascomycin, erythromycin, azithromycin, clarithromycin, clindamycin, lincomycin, dirithromycin, josamycin, spiramycin, diacetyl-midecamycin, tylosin, roxithromycin, ABT-773, telithromycin, leucomycins, and lincosamide.

- 26. (CURRENTLY AMENDED) An article comprising an implantable ocular replacement lens containing at least one macrolide antibiotic or mycophenolic acid at a concentration ranging from about 20 µg/ml to about 2000 µg/ml.
- 27. (ORIGINAL) The article of claim 26 wherein the antibiotic or mycophenolic acid is in a solution in which the lens is contained.
- 28. (ORIGINAL) The article of claim 26 wherein the lens is a porous hydrogel and the antibiotic or mycophenolic acid is within the pores of the hydrogel lens.
- 29. (ORIGINAL) The article of claim 26 wherein the antibiotic or my cophenolic acid is in a coating on at least one lens surface.
- 30. (ORIGINAL) The article of claim 26 wherein the lens is implanted in a lens capsule and the implanted lens releases the antibiotic or mycophenolic acid in the lens capsule.
- 31. (ORIGINAL) The article of claim 26 wherein the macrolide antibiotic is at least one of tacrolimus, cyclosporine, sirolimus, everolimus, ascomycin, erythromycin, azithromycin, clarithromycin, clindamycin, lincomycin, dirithromycin, josamycin, spiramycin, diacetyl-midecamycin, tylosin, roxithromycin, ABT-773, telithromycin, leucomycins, and lincosamide.

32. (CURRENTLY AMENDED) An article comprising an implantable ocular lens in an epthalmically ophthalmically acceptable medium, the medium further comprising an effective anti-cell proliferative or anti-cell migratory concentration of at least one macrolide antibiotic or mycophenolic acid ranging from about 20 μg/ml to about 2000 μg/ml.

33. (CANCELED)

- 34. (ORIGINAL) The article of claim 32 wherein the concentration is in the range between about 200 μg/ml to about 2000 μg/ml.
- 35. (ORIGINAL) The article of claim 32 wherein the concentration is in the range between about 20 μg/ml to about 200 μg/ml.
- 36. (ORIGINAL) The article of claim 32 wherein the macrolide antibiotic is at least one of tacrolimus, cyclosporine, sirolimus, everolimus, ascomycin, erythromycin, azithromycin, clarithromycin, clindamycin, lincomycin, dirithromycin, josamycin, spiramycin, diacetyl-midecamycin, tylosin, roxithromycin, ABT-773, telithromycin, leucomycins, and lincosamide.

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